

IN THE CLAIMS

Please replace all prior versions and claims listings with the following claims listing

Claims Listing:

1-2. (cancel)

3. (previously presented) A method of monitoring for cancer in a biological specimen containing DNA from cells suspected of being cancerous and having PAX5 β gene-specific promoter methylation comprising the steps of:

subjecting DNA to bisulfite modification;

expanding the number of copies of at least a portion of the PAX5 β gene by a polymerase chain reaction to amplify the portion of the PAX5 β gene where the promoter methylation resides, thereby generating an amplification product; and

using an aliquot of the amplification product generated by the first polymerase chain reaction in a second, methylation-specific, polymerase chain reaction at a temperature of annealing that exceeds the melting temperature of the second primer set to amplify a portion of the gene's CpG island where the promoter methylation resides and detect the presence of inactivation of the PAX5 β gene.

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4-6, (cancel)

7. (currently amended) A method of monitoring for cancer in a biological specimen containing DNA from cells suspected of being cancerous and having PAX5 a gene-specific promoter methylation comprising the steps of:

subjecting DNA to **hisulfite** modification;

expanding the number of copies of at least a portion of the PAX5 α gene by using a polymerase chain reaction to amplify a portion of the PAX5 α gene where the promoter methylation resides, thereby generating an amplification product; and

using an aliquot of the amplification product generated by the first polymerase chain reaction in a second, methylation-specific, polymerase chain reaction at a temperature of annealing that exceeds the melting temperature of the second primer set to amplify a portion of the gene's CpG island where the promoter methylation resides and detect the presence of inactivation of the PAX5 α gene.

8-9. (cancel)

10. (previously presented) The method of claim 3 wherein the step of expanding at least a portion of the PAX5 β gene comprises amplifying a 328 base pair fragment with a primer set comprising:

Reverse 5' caaaaaatccccaaaccacccaaaaacc

11. (previously presented) The method of claim 3 wherein the biological sample from which the DNA is obtained is selected from tissue, plasma, ejaculate, cerebrospinal fluid, serum, mammary duct fluid, urine, fecal stool, and sputum.

12. (previously presented) The method of claim 7 wherein the step of expanding at least a portion of the PAX5 α gene comprises amplifying a 389 base pair fragment with a primer set comprising :

Forward 5' gggtttgtatggagatgtatagg

Reverse 5' caacatcacaaaatatccccaaacac

13. (previously presented) The method of claim 7 wherein the biological sample from which the DNA is obtained is selected from tissue, plasma, ejaculate, cerebrospinal fluid, serum, mammary duct fluid, urine, fecal stool, and sputum.